AMENDMENTS TO THE CLAIMS

- 1. (currently amended) A pharmaceutical composition of matter in the form of a solution concentrate compromizing—comprising a cyclosporin dissolved in DMSO dimethyl sulfoxide (DMSO) wherein the concentration of cyclosporin is at least 0.1% by weight of the total composition.
- 2. (currently amended) A composition as in claims claim 1 wherein the cyclosporin is cyclosporin A.
- 3. (currently amended) A method for administering cyclosporin into cerebrospinal fluid spaces, including intraventricular and intrathecal, in a patient, the improvement which compromises:

providing cyclosporin dissolved in DMSO in a pharmaceutically acceptable carrier, and administering said cyclosporin and DMSO solution by injection into the cerebrospinal fluid spaces to said patient.

4. (original) A method for administering cyclosporin by injection including intra-ocular, intravestibular, into or adjacent to the brain, or spinal cord into a patient, the improvement which compromises: providing cyclosporin dissolved in DMSO in a

Appl. No. 09/674,092

pharmaceutically acceptable carrier, and administering said cyclosporin and DMSO solution by injection intra-ocular, intravestibular, into or adjacent to the brain, or spinal cord to said patient.

- 5. (original) A method for administering cyclosporin by injection including intravenous, intra-arterial orinto a intraparenchymal, patient, the improvement which compromises: providing cyclosporin dissolved in DMSO pharmaceutically acceptable carrier, and administering cyclosporin and DMSO solution by injection into intravenous, intraarterial or intraparenchymal spaces to said patient.
- 6. (original) A method for administering cyclosporin orally, rectally, nasally or dermally to a patient, the improvement which compromises: providing the cyclosporin dissolved in DMSO in a pharmaceutically acceptable carrier, and administering said cyclosporin and DMSO solution orally, rectally, nasally or dermally to said patient.
- 7. (previously amended) The method of claim 3 wherein the cyclosporin is cyclosporin A, or functional derivatives, metabolites, variants or salts thereof.

- 8. (currently amended) An article of manufacture comprising packaging material and pharmaceutical agent is therapeutically effective for reducing or preventing neuronal damage and for causing immunosuppression when administered in a therapeutically effective quantity, and wherein the packaging material comprises a label which indicates that the pharmaceutical agent can be used for preventing neuronal damage reducing or and for immunosuppression, and wherein said pharmaceutical agent comprises DMSO and a cyclosporin such as cyclosporin A or a compound of the class of cyclosporins, or functional derivatives, metabolites, variants or salts of them thereof, or a combination of the before said, either alone or in admixture with diluents, or additives.
- 9. (new) The article of manufacture according to claim 8, wherein the cyclosporin is selected from the group consisting of cyclosporin A and a compound of the class of cyclosporins.
- 10. (new) The method according to claim 3 wherein the administering cyclosporin into cerebrospinal fluid spaces is intraventricular and intrathecal.